

**K103176 FREQUENCER**Jan 26, 2011  
90 days to decisionK103176 · Product code: **BYI** · Anesthesiology  
Source: <https://www.510kdatabase.net/k103176/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percussor, Powered-electric (BYI)
Date received	Oct 28, 2010
Decision date	Jan 26, 2011
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dymedso, Inc.</b>
Location	Springfield, VT, US
Contact	Jean Bigoney
510(k) history	3 submissions · 3 cleared · 2007-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103176/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 24, 2026